

09/979453
L/cook 1/11/05

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(FILE 'HOME' ENTERED AT 11:32:09 ON 11 JAN 2005)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, CANCERLIT, JAPIO' ENTERED AT
11:32:37 ON 11 JAN 2005

L1 129 S (FLUID FLOW CHANNEL)
L2 0 S L1 AND (MULTIPLE DETECT?)
L3 14 S L1 AND DETECT?
L4 13 DUPLICATE REMOVE L3 (1 DUPLICATE REMOVED)
L5 0 S L4 AND MULTIPL?
L6 1 S L4 AND PLURALI?
L7 6972 S MICROFLUIDIC?
L8 5 S L7 AND (MULTIPLE DETECT?)
L9 3 DUPLICATE REMOVE L8 (2 DUPLICATES REMOVED)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, CANCERLIT, JAPIO' ENTERED AT
11:55:00 ON 11 JAN 2005

L10 6972 S MICROFLUIDIC?
L11 0 S L10 AND (DUAL DETECTOR?)
L12 352 S L10 AND DETECTOR?
L13 14 S L12 AND VELOCIT?
L14 11 DUPLICATE REMOVE L13 (3 DUPLICATES REMOVED)

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L7 6972 S MICROFLUIDIC?
L8 5 S L7 AND (MULTIPLE DETECT?)
L9 3 DUPLICATE REMOVE L8 (2 DUPLICATES REMOVED)

=>

ANSWER 3 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1

AN 2001:162337 CAPLUS

ED Entered STN: 08 Mar 2001

TI Velocity measurement of particles flowing in a **microfluidic** chip using Shah convolution Fourier transform detection

AU Kwok, Yien C.; Jeffery, Nicholas T.; Manz, Andreas

CS Department of Chemistry, AstraZeneca/SmithKline Beecham Centre for Analytical Sciences Imperial College of Science Technology and Medicine, London, SW7 2AY, UK

SO Analytical Chemistry ~~(2001)~~ 73(8), 1748-1753

CODEN: ANCHAM; ISSN: 0003-2700

PB American Chemical Society

DT Journal

LA English

AB A noninvasive radiative technique, based on Shah convolution Fourier transform detection, for velocity measurement of particles in fluid flows in a **microfluidic** chip, is presented. It boasts a simpler instrumental setup and optical alignment than existing measurement methods and a wide dynamic range of velocities measurable. A glass-PDMS microchip with a layer of patterned Cr to provide **multiple detection** windows which are 40 μm wide and 70 μm apart is employed. The velocities of fluorescent microspheres, which were electrokinetically driven in the channel of the **microfluidic** chip, were determined. The effects of increasing the number of detection

windows

and sampling period were investigated. This technique could have wide applications, ranging from the determination of the velocity of particles in pressure-driven flow to the measurement of electrophoretic mobilities of single biol. cells.

date no good

ANSWER 3 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1

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and sampling period were investigated. This technique could have wide
applications, ranging from the determination of the velocity of particles in
pressure-driven flow to the measurement of electrophoretic mobilities of
single biol. cells.

ANSWER 1 OF 3 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation. on STN
AN 2003:227200 BIOSIS
DN PREV200300227200
TI Ultra high throughput **microfluidic** analytical systems and
methods.
AU Kopf-Sill, Anne R. [Inventor, Reprint Author]; Chow, Andrea W. [Inventor];
Jann, Peter C. [Inventor]; Jensen, Morten J. [Inventor]; Spaid, Michael
[Inventor]; Kennedy, Colin B. [Inventor]; Kennedy, Michael J. [Inventor]
CS Santa Clara, CA, USA
ASSIGNEE: Caliper Technologies Corp.
PI US 6547941 April 15, 2003
SO Official Gazette of the United States Patent and Trademark Office Patents,
(Apr 15 2003) Vol. 1269, No. 3. <http://www.uspto.gov/web/menu/patdata.html>
. e-file.
ISSN: 0098-1133 (ISSN print).
DT Patent
LA English
ED Entered STN: 7 May 2003
Last Updated on STN: 7 May 2003
AB Analytical systems and methods that use a modular interface structure for
providing an interface between a sample substrate and an analytical unit,
where the analytical unit typically has a particular interface arrangement
for implementing various analytical and control functions. Using a number
of variants for each module of the modular interface structure
advantageously provides cost effective and efficient ways to perform
numerous tests using a particular substrate or class of substrates with a
particular analytical and control systems interface arrangement. Improved
optical illumination and detection system for simultaneously analyzing
reactions or conditions in multiple parallel microchannels are also
provided. Increased throughput and improved emissions detection is
provided by the present invention by simultaneously illuminating multiple
parallel microchannels at a non-normal incidence using an excitation beam
including multiple excitation frequencies, and simultaneously detecting
emissions from the substances in the microchannels in a direction normal
to the substrate using a detection module with **multiple**
detectors.
NCL 204452000
CC Biochemistry studies - General 10060
IT Major Concepts
Chemistry; Methods and Techniques
IT Methods & Equipment
high throughput **microfluidic** analytical systems: laboratory
equipment; ultra high throughput **microfluidic** analytical
methods: laboratory techniques

✓ pull patent.
filing date. -

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:174333 CAPLUS
 DN 138:201292
 ED Entered STN: 07 Mar 2003
 TI Analysis using a distributed sample
 IN Matson, Robert S.
 PA USA
 SO U.S. Pat. Appl. Publ., 11 pp.
 CODEN: USXXCO
 DT Patent
 LA English
 IC ICM C12Q001-68
 ICS G01N033-53; G01N033-542
 NCL 435006000; 435007900
 CC 9-1 (Biochemical Methods)
 Section cross-reference(s): 3, 15

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003044799	A1	20030306	US 2001-945145	20010831
PRAI	US 2001-945145		20010831		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
US 2003044799	ICM	C12Q001-68
	ICS	G01N033-53; G01N033-542
	NCL	435006000; 435007900

AB The present invention is directed to the production of a sample microarray for use in detecting one or more target biopolymers in the sample. The sample microarray of this invention is formed by distributing equivalent amts. of a single sample at **discrete**, spatially defined **locations** on a substrate. Each site in the microarray, thus, has the same composition of target biopolymers. The microarray is then interrogated by one or more probes specific for one or more the target biopolymers.

ST microarray detecting target biopolymer

IT Functional groups

(Alkanethiol; anal. using distributed sample)

IT Printing (impact)

(Capillary quill contact; anal. using distributed sample)

IT Fluoropolymers, uses

RL: DEV (Device component use); USES (Uses)

(Carboxylated; anal. using distributed sample)

IT Polymers, uses

RL: DEV (Device component use); USES (Uses)

(Crosslinked; anal. using distributed sample)

IT Adhesives

(Die-cut; anal. using distributed sample)

IT Antibodies and Immunoglobulins

RL: ANT (Analyte); ANST (Analytical study)

(IgG; anal. using distributed sample)

IT Printing (impact)

(Microfluidic-based; anal. using distributed sample)

IT Materials

(Nonporous metallic; anal. using distributed sample)

IT Apparatus

(Planar; anal. using distributed sample)

IT Apparatus

(Radio frequency transmitters; anal. using distributed sample)

IT Biochemical molecules

(Radioactive-labeled; anal. using distributed sample)

IT Molecules

(Radioluminescent; anal. using distributed sample)

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:174333 CAPLUS
 DN 138:201292
 ED Entered STN: 07 Mar 2003
 TI Analysis using a distributed sample
 IN Matson, Robert S.
 PA USA
 SO U.S. Pat. Appl. Publ., 11 pp.
 CODEN: USXXCO
 DT Patent
 LA English
 IC ICM C12Q001-68
 ICS G01N033-53; G01N033-542
 NCL 435006000; 435007900
 CC 9-1 (Biochemical Methods)
 Section cross-reference(s): 3, 15

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PRAI	US 2001-945145		20010831		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
US 2003044799	ICM	C12Q001-68
	ICS	G01N033-53; G01N033-542
	NCL	435006000; 435007900

AB The present invention is directed to the production of a sample microarray for use in detecting one or more target biopolymers in the sample. The sample microarray of this invention is formed by distributing equivalent amts. of a single sample at **discrete**, spatially defined **locations** on a substrate. Each site in the microarray, thus, has the same composition of target biopolymers. The microarray is then interrogated by one or more probes specific for one or more the target biopolymers.

ST microarray detecting target biopolymer

IT Functional groups

(Alkanethiol; anal. using distributed sample)

IT Printing (impact)

(Capillary quill contact; anal. using distributed sample)

IT Fluoropolymers, uses

RL: DEV (Device component use); USES (Uses)

(Carboxylated; anal. using distributed sample)

IT Polymers, uses

RL: DEV (Device component use); USES (Uses)

(Crosslinked; anal. using distributed sample)

IT Adhesives

(Die-cut; anal. using distributed sample)

IT Antibodies and Immunoglobulins

RL: ANT (Analyte); ANST (Analytical study)

(IgG; anal. using distributed sample)

IT Printing (impact)

(Microfluidic-based; anal. using distributed sample)

IT Materials

(Nonporous metallic; anal. using distributed sample)

IT Apparatus

(Planar; anal. using distributed sample)

IT Apparatus

(Radio frequency transmitters; anal. using distributed sample)

IT Biochemical molecules

(Radioactive-labeled; anal. using distributed sample)

IT Molecules

(Radioluminescent; anal. using distributed sample)

IT Printing (impact)
(Solid pin; anal. using distributed sample)

IT Materials
(Surface modified; anal. using distributed sample)

IT Materials
(Surface-modified; anal. using distributed sample)

IT Acid halides
RL: ANT (Analyte); ANST (Analytical study)
(acid fluorides; anal. using distributed sample)

IT DNA
RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
(amplified; anal. using distributed sample)

IT Absorption
Adsorption
Amino group
Animal tissue
Bar code labels
Carboxyl group
Cell
Ceramics
Chemiluminescent substances
Composition
Concentration (condition)
DNA microarray technology
Drugs
Dyes
Filaments
Films
Flow
Fluids
Fluorescent indicators
Foams
Functional groups
Gels
Heating
Human
Hydroxyl group
Immobilization, molecular or cellular
Ink-jet printing
Magnetic particles
Membranes, nonbiological
Microarray technology
Microtiter plates
Nucleic acid hybridization
Particles
Plates
Protein microarray technology
Quantum dot devices
Samples
Solenoids
Surface area
Threads
Wells
Wetting
(anal. using distributed sample)

IT Biopolymers
Nucleic acids
Organic compounds, analysis
Proteins
RL: ANT (Analyte); ANST (Analytical study)
(anal. using distributed sample)

IT Printing (impact)
 (Solid pin; anal. using distributed sample)

IT Materials
 (Surface modified; anal. using distributed sample)

IT Materials
 (Surface-modified; anal. using distributed sample)

IT Acid halides
 RL: ANT (Analyte); ANST (Analytical study)
 (acid fluorides; anal. using distributed sample)

IT DNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (amplified; anal. using distributed sample)

IT Absorption
 Adsorption
 Amino group
 Animal tissue
 Bar code labels
 Carboxyl group
 Cell
 Ceramics
 Chemiluminescent substances
 Composition
 Concentration (condition)
 DNA microarray technology
 Drugs
 Dyes
 Filaments
 Films
 Flow
 Fluids
 Fluorescent indicators
 Foams
 Functional groups
 Gels
 Heating
 Human
 Hydroxyl group
 Immobilization, molecular or cellular
 Ink-jet printing
 Magnetic particles
 Membranes, nonbiological
 Microarray technology
 Microtiter plates
 Nucleic acid hybridization
 Particles
 Plates
 Protein microarray technology
 Quantum dot devices
 Samples
 Solenoids
 Surface area
 Threads
 Wells
 Wetting
 (anal. using distributed sample)

IT Biopolymers
 Nucleic acids
 Organic compounds, analysis
 Proteins
 RL: ANT (Analyte); ANST (Analytical study)
 (anal. using distributed sample)

IT Carbohydrates, analysis
 Peptide nucleic acids
 Polynucleotides
 Receptors
 cDNA
 mRNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (anal. using distributed sample)

IT Antibodies and Immunoglobulins
 Antigens
 Coordination compounds
 Enzymes, uses
 Haptens
 Ligands
 Probes (nucleic acid)
 RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (anal. using distributed sample)

IT Epoxides
 Esters, uses
 Glass, uses
 Polyamides, uses
 RL: DEV (Device component use); USES (Uses)
 (anal. using distributed sample)

IT Spheres
 (beads, Dye-labeled; anal. using distributed sample)

IT Spheres
 (beads; anal. using distributed sample)

IT Bond
 (covalent; anal. using distributed sample)

IT DNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (double-stranded; anal. using distributed sample)

IT Antibodies and Immunoglobulins
 RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (fragments; anal. using distributed sample)

IT Standard substances, analytical
 (internal; anal. using distributed sample)

IT Porous materials
 (metallic; anal. using distributed sample)

IT Peptides, analysis
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (polypeptides; anal. using distributed sample)

IT Printing (nonimpact)
 (silk-screen; anal. using distributed sample)

IT DNA
 RNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (single-stranded; anal. using distributed sample)

IT Laboratory ware
 (slides; anal. using distributed sample)

IT Containers
 (troughs; anal. using distributed sample)

IT 58-85-5, Biotin
 RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (anal. using distributed sample)

IT Carbohydrates, analysis
 Peptide nucleic acids
 Polynucleotides
 Receptors
 cDNA
 mRNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (anal. using distributed sample)

IT Antibodies and Immunoglobulins
 Antigens
 Coordination compounds
 Enzymes, uses
 Haptens
 Ligands
 Probes (nucleic acid)
 RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (anal. using distributed sample)

IT Epoxides
 Esters, uses
 Glass, uses
 Polyamides, uses
 RL: DEV (Device component use); USES (Uses)
 (anal. using distributed sample)

IT Spheres
 (beads, Dye-labeled; anal. using distributed sample)

IT Spheres
 (beads; anal. using distributed sample)

IT Bond
 (covalent; anal. using distributed sample)

IT DNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (double-stranded; anal. using distributed sample)

IT Antibodies and Immunoglobulins
 RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (fragments; anal. using distributed sample)

IT Standard substances, analytical
 (internal; anal. using distributed sample)

IT Porous materials
 (metallic; anal. using distributed sample)

IT Peptides, analysis
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (polypeptides; anal. using distributed sample)

IT Printing (nonimpact)
 (silk-screen; anal. using distributed sample)

IT DNA
 RNA
 RL: ANT (Analyte); ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (single-stranded; anal. using distributed sample)

IT Laboratory ware
 (slides; anal. using distributed sample)

IT Containers
 (troughs; anal. using distributed sample)

IT 58-85-5, Biotin
 RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 (anal. using distributed sample)

IT 7440-57-5, Gold, uses 7631-86-9, Silica, uses 9002-88-4, Polyethylene
9003-01-4, Polyacrylic acid 9003-07-0, Polypropylene 9003-53-6,
Polystyrene 9004-70-0, Nitrocellulose 24937-79-9D, Polyvinylidene
fluoride, Carboxylated
RL: DEV (Device component use); USES (Uses)
(anal. using distributed sample)

IT 64-17-5, Ethanol, uses 64-19-7, Acetic acid, uses 67-56-1, Methanol,
uses 67-63-0, Isopropanol, uses 78-92-2, 2-Butanol 9042-14-2,
Dextran sulfate
RL: NUU (Other use, unclassified); USES (Uses)
(anal. using distributed sample)

IT 7440-57-5, Gold, uses 7631-86-9, Silica, uses 9002-88-4, Polyethylene
9003-01-4, Polyacrylic acid 9003-07-0, Polypropylene 9003-53-6,
Polystyrene 9004-70-0, Nitrocellulose 24937-79-9D, Polyvinylidene
fluoride, Carboxylated
RL: DEV (Device component use); USES (Uses)
(anal. using distributed sample)

IT 64-17-5, Ethanol, uses 64-19-7, Acetic acid, uses 67-56-1, Methanol,
uses 67-63-0, Isopropanol, uses 78-92-2, 2-Butanol 9042-14-2,
Dextran sulfate
RL: NUU (Other use, unclassified); USES (Uses)
(anal. using distributed sample)

AN 1999:183707 CAPLUS
DN 130:317031
ED Entered STN: 22 Mar 1999
TI MEMS based micro-fluidic system for chromatographic analysis of liquid samples
AU Golubovic, Nevenka C.; Kang, Qinghua; Henderson, H. Thurman; Pinto, Neville
CS Center for Microelectronic Sensors, CMSM, Cincinnati, OH, 45221-0030, USA
SO Proceedings of SPIE-The International Society for Optical Engineering (1998), 3515(Microfluidic Devices and Systems), 86-93
CODEN: PSISDG; ISSN: 0277-786X
PB SPIE-The International Society for Optical Engineering
DT Journal
LA English
CC 66-4 (Surface Chemistry and Colloids)
Section cross-reference(s): 47, 79, 80
AB A complete micro-chromatog. system has been designed on a (110) silicon chip and the column-detector sub-system has been demonstrated. This micro-configuration allows the active surface-to-cross sectional area to be maximized, consistent with fabrication and pressure drop issues. A separation column was designed as an array of parallel channels anisotropically etched in (110) silicon to reduce pressure drop and to provide a necessary large surface area at a short length. Sensing was done by use of integrated impedance electrodes, with the detector cell volume less than 1 nl, although integrated optical detection has also been initiated. The response time is improved by about two orders of magnitude (relative to traditional systems) and simultaneous multiple anal. capability is realized with this design. Fabrication of multiple impedance **detectors** at different **locations** along the length of a micro-channel will enable monitoring of the separation in progress. Although the present work supports only a linear column configuration, a serpentine version would consume only about one square millimeter of a chip area, thus further minimizing the device.
ST MEMS **microfluidic** system open tubular liq chromatog
IT Liquid chromatography
Sensors
(MEMS based micro-fluidic system for chromatog. anal. of liquid samples)
IT 7440-21-3, Silicon, processes
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
(miniature liquid chromatog. device fabricated on a silicon chip)
RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) Guiochon, G; Analytical Chemistry 1981, V53, P1318 CAPLUS
(2) Ishii, D; Advances in Chromatography 1983, V21, P131 CAPLUS
(3) Manz, A; Sens Actuators 1990, VB1, P249 CAPLUS
(4) Ocvirk, G; Proc Transducers '95 1995, P756
(5) Reston, R; IEEE J Microelectromech Syst 1994, V3(4), P134 CAPLUS
(6) Reston, R; IEEE J Microelectromech Syst 1994, V3(4), P147
(7) Terry, S; IEEE Trans Electron Devices 1979, VED-26(12), P1880 CAPLUS
(8) Tijssen, R; J of Chromatography 1981, V218, P137 CAPLUS